Docket No: 966927.00005 (0273-0005) Appl. No: 10/026,911

In the Claims:

Please cancel claim 7 and amend claim 1 as shown in the following listing of the entire claims in the Application.

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1. (currently amended) A pharmaceutical composition containing a peptide wherein all

amino acids of the peptide except one are identical to the amino acids of an amino acid

sequence which is part of an allergenic protein amino acid sequence and the allergenic

protein is the birch pollen allergen Bet v 1 and a pharmaceutically acceptable carrier or

diluent, wherein the peptide:

a) has a length of 8 to 50 amino acids; [[and]]

b) has at least three preferably consecutive amino acids identical to at least

three solvent-exposed amino acids of an allergenic protein which appear in close

vicinity on the molecular surface of the allergenic protein; and

the peptide leads upon administration to the production of protective IgG

antibodies which react with the protein from which the peptide is derived.

2. (previously presented) A pharmaceutical composition according to claim 1, wherein

said at least three solvent-exposed amino acids appear on the molecular surface of the

allergenic protein within a surface patch of approximately 500 square Angström.

3. (previously presented) A pharmaceutical composition according to claim 1, wherein at

least five consecutive amino acids of the peptide are identical to at least five consecutive

solvent-exposed amino acids of the allergenic protein.

4. (previously presented) A pharmaceutical composition according to claim 1, further

containing an adjuvant.

5. (canceled)

6. (currently amended) A pharmaceutical composition according to claim 1, wherein the

one amino acid which deviates from the amino acid sequence of the allergenic protein is

the N-terminal or C-terminal amino acid of the peptide amino acid sequence.

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7. (canceled)

8. (canceled)

9. (previously presented) A pharmaceutical composition according to claim 1, wherein

the peptide amino acid sequence comprises at least the N-terminal or C-terminal five

amino acids of the allergenic protein amino acid sequence.

10 - 13. (canceled).

14. (withdrawn) A method for preparing a pharmaceutical composition comprising:

a) determining which amino acids of a given allergenic protein are solvent-

exposed on the surface of the allergenic protein;

b) preparing a peptide having a length of 8 to 50 amino acids, wherein at

least three preferably consecutive amino acids of the peptide are identical to at

least three solvent-exposed amino acids of the allergenic protein which appear in

close vicinity on the molecular surface of the allergenic protein; and

c) optionally admixing the peptide with a pharmaceutically acceptable carrier

or diluent.

15. (withdrawn) A method according to claim 14, wherein said at least three solvent-

exposed amino acids appear on the molecular surface of the allergenic protein within a

surface patch of approximately 500 square Angström.

16. (withdrawn) A method according to claim 14, wherein at least

five consecutive amino acids of the peptide are identical to at least five consecutive

solvent-exposed amino acids of the allergenic protein.

17. (withdrawn) A method according to claim 14, further comprising adding an adjuvant.

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18. (withdrawn) A method according to claim 14, wherein all amino acids of the peptide

except one are identical to the amino acids of an amino acid sequence which is part of the

allergenic protein amino acid sequence.

19. (withdrawn) A method according to claim 18, wherein the one amino acid which

deviates from the amino acid sequence of the allergenic protein is the N-terminal or C-

terminal amino acid of the peptide amino acid sequence.

20. (withdrawn) A method according to claim 14, wherein the amino acid sequence of

the peptide is identical to an amino acid sequence which is part of the allergenic protein

amino acid sequence.

21. (withdrawn) A method according to claim 14, wherein the allergenic protein is the

birch pollen allergen Bet v 1.

22. (withdrawn) A method according to claim 14, wherein the peptide amino acid

sequence comprises at least the N-terminal or C-terminal five amino acids of the

allergenic protein amino acid sequence.

23. (withdrawn) A method according to claim 14, wherein the solvent-exposed amino

acids of the allergenic protein are determined by determining the hydrophilicity profile of

the allergenic protein.

24. (withdrawn) A method according to claim 14, wherein the solvent-exposed amino

acids of the allergenic protein are determined from the three-dimensional structure of the

allergenic protein.

25 - 27 (canceled).

28. (withdrawn) A method for treating an allergic disease, comprising: administering

to a patient in need thereof the pharmaceutical composition of claim 1.

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29. (withdrawn) A method according to claim 28, wherein the at least three solvent-exposed amino acids appear on the molecular surface of the allergenic protein within a surface patch of approximately 500 square Angström.

30. (withdrawn) A method according to claim 28, wherein at least five consecutive amino acids of the peptide are identical to at least five consecutive solvent-exposed amino acids of the allergenic protein.

31. (withdrawn) A method according to claim 28, wherein the peptide, upon administration, is capable of inducing IgG antibodies which react with the allergenic protein.

32. (withdrawn) A method according to claim 31, wherein the induced IgG antibodies can reduce or prevent binding of IgE antibodies to the allergenic protein.

33. (withdrawn) A method according to claim 28, wherein the peptide, upon administration, does not induce a significant IgE response.